EAST Search History

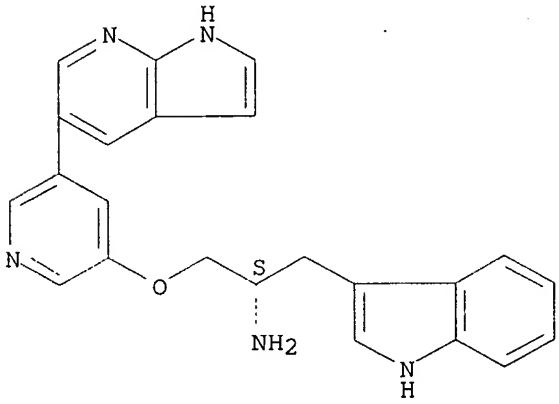
Ref #	Hits	Search Query	DBs	Default Operator	Plurals	Time Stamp
L1	784	(546/113).CCLS.	US-PGPUB; USPAT	OR	OFF	2006/11/07 17:26
L2	1514	(514/300).CCLS.	US-PGPUB; USPAT	OR	OFF	2006/11/07 17:26
L3	4067	pyrrolo	US-PGPUB; USPAT	OR	ON	2006/11/07 17:26
L4	208	l1 and l3	US-PGPUB; USPAT	OR	ON	2006/11/07 17:26
L5	210	l2 and l3	US-PGPUB; USPAT	OR	ON	2006/11/07 17:26
L6	167586	pyridine	US-PGPUB; USPAT	OR	ON	2006/11/07 17:26
L7	187	l4 and l6	US-PGPUB; USPAT	OR	ON	2006/11/07 17:26
L8	189	l5 and l6	US-PGPUB; USPAT	OR	ON	2006/11/07 17:27
L9	125	"[2,3-b]"	US-PGPUB; USPAT	OR	ON	2006/11/07 17:27
L10	9	l7 and l9	US-PGPUB; USPAT	OR	ON	2006/11/07 17:27
L11	7	l8 and l9	US-PGPUB; USPAT	OR	ON	2006/11/07 17:27
L12	4	piotr and graczyk	US-PGPUB; USPAT	OR	ON	2006/11/07 17:28
L13	2	hirotoshi and numata and london	US-PGPUB; USPAT	OR	ON	2006/11/07 17:28
L14	5	gurpreet and bhatia	US-PGPUB; USPAT	OR	ON	2006/11/07 17:29
L15	2	darren and peter and medland	US-PGPUB; USPAT	OR	ON	2006/11/07 17:28

L9 ANSWER 1 OF 8 CAPLUS COPYRIGHT 2006 ACS on STN
ACCESSION NUMBER: 2003:836590 CAPLUS
DOCUMENT NUMBER: 139:323437
TITLE: Preparation of heteroaryls for therapeutic use in pharmaceutical compositions as kinase inhibitors for treatment of hyperproliferative diseases, including cancer
INVENTOR(S): Li, Qun; Woods, Keith W.; Zhu, Gui-Dong; Fischer, John P.; Gong, Jianchun; Li, Tongmei; Gandhi, Virajkumar; Thomas, Sheela A.; Packard, Garrick K.; Song, Xiaohong; Abrams, Jason N.; Diebold, Robert B.; Dinges, Jurgen; Hutchins, Charles W.; Stoll, Vincent S.; Rosenberg, Saul H.; Giranda, Vincent L.
PATENT ASSIGNEE(S): Abbott Laboratories, USA
SOURCE: U.S. Pat. Appl. Publ., 120 pp., which
CODEN: USXXCO
DOCUMENT TYPE: Patent
LANGUAGE: English
FAMILY ACC. NUM. COUNT: 1
PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.		DATE
US 2003199511	A1	20031023	US 2002-317914		20021212 <--
US 6831175	B2	20041214			
PRIORITY APPLN. INFO.:			US 2001-341356P	P	20011213
			US 2001-341474P	P	20011217

OTHER SOURCE(S): MARPAT 139:323437
IT 552326-49-5P 552326-50-8P
RL: PAC (Pharmacological activity); SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)
(preparation of heteroaryls for therapeutic use in pharmaceutical compns. as kinase inhibitors for treatment of hyperproliferative diseases, including cancer)
RN 552326-49-5 CAPLUS
CN 1H-Indole-3-ethanamine, α-[[[5-(1H-pyrrolo[2,3-b]pyridin-5-yl)-3-pyridinyl]oxy]methyl]-, (αS)- (9CI) (CA INDEX NAME)

Absolute stereochemistry.

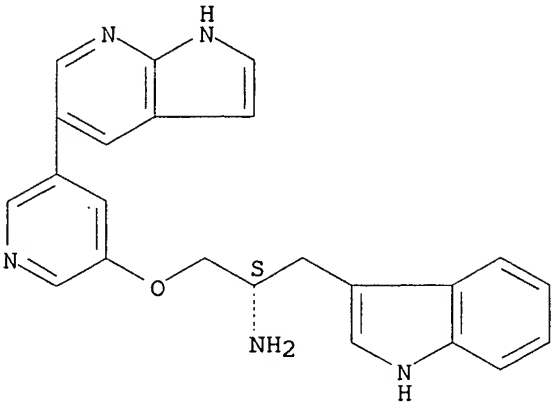


RN 552326-50-8 CAPLUS
CN 1H-Indole-3-ethanamine, α-[[[5-(1H-pyrrolo[2,3-b]pyridin-5-yl)-3-pyridinyl]oxy]methyl]-, (αS)-, trifluoroacetate (9CI) (CA INDEX NAME)

CM 1

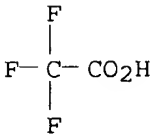
CRN 552326-49-5
CMF C23 H21 N5 O

Absolute stereochemistry.



CM 2

CRN 76-05-1
CMF C2 H F3 O2



REFERENCE COUNT: 64 THERE ARE 64 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L9 ANSWER 2 OF 8 CAPLUS COPYRIGHT 2006 ACS on STN
ACCESSION NUMBER: 2003:796705 CAPLUS
DOCUMENT NUMBER: 139:307750
TITLE: Preparation of 7-azaindoles as inhibitors of c-Jun N-terminal kinases
INVENTOR(S): Graczyk, Piotr; Numata, Hirotoshi; Bhatia, Gurpreet; Medland, Darren Peter
PATENT ASSIGNEE(S): Eisai London Research Laboratories Limited, UK
SOURCE: PCT Int. Appl., 44 pp.
CODEN: PIXXD2
DOCUMENT TYPE: Patent
LANGUAGE: English
FAMILY ACC. NUM. COUNT: 1
PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
WO 2003082869	A1	20031009	WO 2003-GB1115	20030317 <--

W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE, GH,

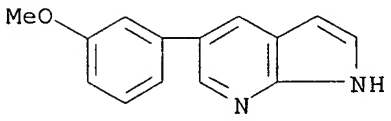
GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR,
LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NI, NO, NZ, OM,
PH, PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, TJ, TM, TN, TR, TT,
TZ, UA, UG, US, UZ, VC, VN, YU, ZA, ZM, ZW
RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ, BY,
KG, KZ, MD, RU, TJ, TM, AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES,
FI, FR, GB, GR, HU, IE, IT, LU, MC, NL, PT, RO, SE, SI, SK, TR,
BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG

CA 2479205	AA	20031009	CA 2003-2479205		20030317 <--
AU 2003214414	A1	20031013	AU 2003-214414		20030317 <--
EP 1490365	A1	20041229	EP 2003-709986		20030317
R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, SI, LT, LV, FI, RO, MK, CY, AL, TR, BG, CZ, EE, HU, SK					
CN 1649869	A	20050803	CN 2003-809569		20030317
JP 2005534619	T2	20051117	JP 2003-580334		20030317
PRIORITY APPLN. INFO.:			GB 2002-7488	A	20020328
			GB 2003-400	A	20030108
			WO 2003-GB1115	W	20030317

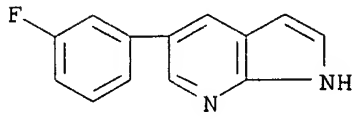
OTHER SOURCE(S): MARPAT 139:307750

IT 344454-28-0P 611204-92-3P 611204-93-4P
611204-94-5P 611204-95-6P 611204-96-7P
611204-97-8P 611204-98-9P 611204-99-0P
611205-00-6P 611205-01-7P 611205-02-8P
611205-03-9P 611205-04-0P 611205-05-1P
611205-06-2P 611205-07-3P 611205-08-4P
611205-09-5P 611205-10-8P 611205-11-9P
611205-12-0P 611205-13-1P 611205-14-2P
611205-15-3P 611205-16-4P 611205-17-5P
611205-18-6P 611205-19-7P 611205-20-0P
611205-21-1P 611205-22-2P 611205-23-3P
611205-24-4P 611205-25-5P 611205-26-6P
611205-27-7P 611205-28-8P 611205-29-9P
611205-30-2P 611205-31-3P 611205-32-4P
611205-33-5P 611205-34-6P 611205-35-7P
611205-36-8P 611205-37-9P 611205-38-0P
611205-39-1P 611205-40-4P 611205-41-5P
611205-42-6P 611205-43-7P
RL: PAC (Pharmacological activity); SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)
(drug candidate; preparation of azaindoles as inhibitors of c-jun N-terminal kinases)

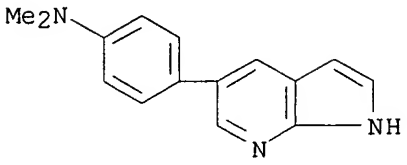
RN 344454-28-0 CAPLUS
CN 1H-Pyrrolo[2,3-b]pyridine, 5-(3-methoxyphenyl)- (9CI) (CA INDEX NAME)



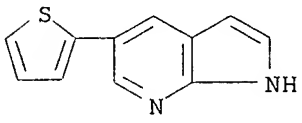
RN 611204-92-3 CAPLUS
CN 1H-Pyrrolo[2,3-b]pyridine, 5-(3-fluorophenyl)- (9CI) (CA INDEX NAME)



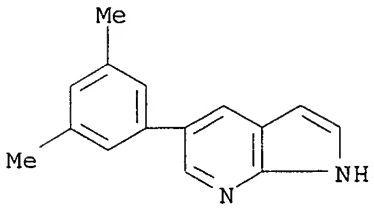
RN 611204-93-4 CAPLUS
CN Benzenamine, N,N-dimethyl-4-(1H-pyrrolo[2,3-b]pyridin-5-yl)- (9CI) (CA INDEX NAME)



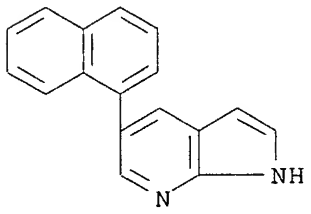
RN 611204-94-5 CAPLUS
CN 1H-Pyrrolo[2,3-b]pyridine, 5-(2-thienyl)- (9CI) (CA INDEX NAME)



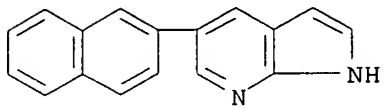
RN 611204-95-6 CAPLUS
CN 1H-Pyrrolo[2,3-b]pyridine, 5-(3,5-dimethylphenyl)- (9CI) (CA INDEX NAME)



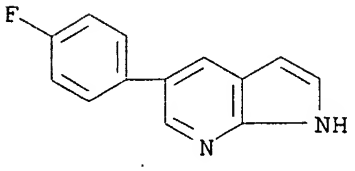
RN 611204-96-7 CAPLUS
CN 1H-Pyrrolo[2,3-b]pyridine, 5-(1-naphthalenyl)- (9CI) (CA INDEX NAME)



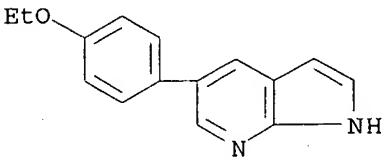
RN 611204-97-8 CAPLUS
CN 1H-Pyrrolo[2,3-b]pyridine, 5-(2-naphthalenyl)- (9CI) (CA INDEX NAME)



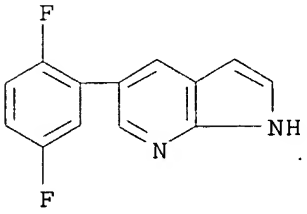
RN 611204-98-9 CAPLUS
CN 1H-Pyrrolo[2,3-b]pyridine, 5-(4-fluorophenyl)- (9CI) (CA INDEX NAME)



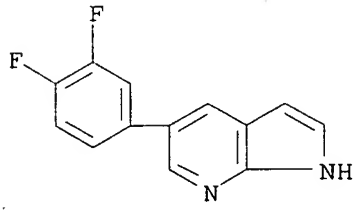
RN 611204-99-0 CAPLUS
CN 1H-Pyrrolo[2,3-b]pyridine, 5-(4-ethoxyphenyl)- (9CI) (CA INDEX NAME)



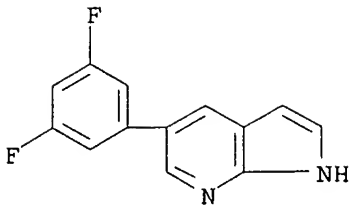
RN 611205-00-6 CAPLUS
CN 1H-Pyrrolo[2,3-b]pyridine, 5-(2,5-difluorophenyl)- (9CI) (CA INDEX NAME)



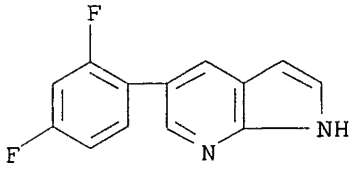
RN 611205-01-7 CAPLUS
CN 1H-Pyrrolo[2,3-b]pyridine, 5-(3,4-difluorophenyl)- (9CI) (CA INDEX NAME)



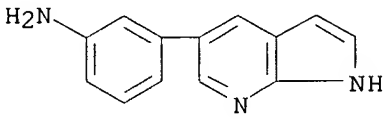
RN 611205-02-8 CAPLUS
CN 1H-Pyrrolo[2,3-b]pyridine, 5-(3,5-difluorophenyl)- (9CI) (CA INDEX NAME)



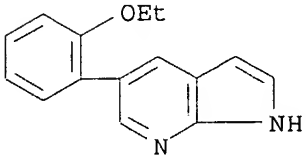
RN 611205-03-9 CAPLUS
CN 1H-Pyrrolo[2,3-b]pyridine, 5-(2,4-difluorophenyl)- (9CI) (CA INDEX NAME)



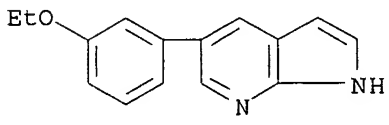
RN 611205-04-0 CAPLUS
CN Benzenamine, 3-(1H-pyrrolo[2,3-b]pyridin-5-yl)- (9CI) (CA INDEX NAME)



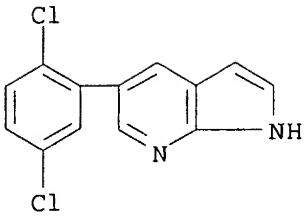
RN 611205-05-1 CAPLUS
CN 1H-Pyrrolo[2,3-b]pyridine, 5-(2-ethoxyphenyl)- (9CI) (CA INDEX NAME)



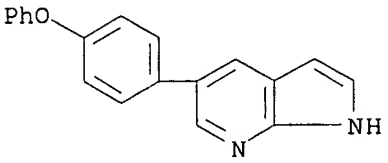
RN 611205-06-2 CAPLUS
CN 1H-Pyrrolo[2,3-b]pyridine, 5-(3-ethoxyphenyl)- (9CI) (CA INDEX NAME)



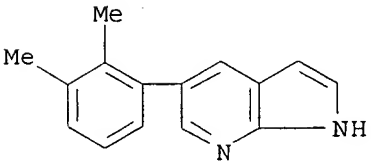
RN 611205-07-3 CAPLUS
CN 1H-Pyrrolo[2,3-b]pyridine, 5-(2,5-dichlorophenyl)- (9CI) (CA INDEX NAME)



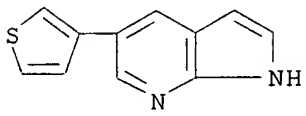
RN 611205-08-4 CAPLUS
CN 1H-Pyrrolo[2,3-b]pyridine, 5-(4-phenoxyphenyl)- (9CI) (CA INDEX NAME)



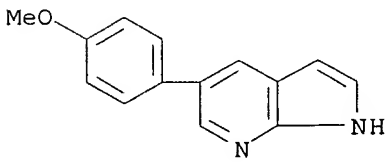
RN 611205-09-5 CAPLUS
CN 1H-Pyrrolo[2,3-b]pyridine, 5-(2,3-dimethylphenyl)- (9CI) (CA INDEX NAME)



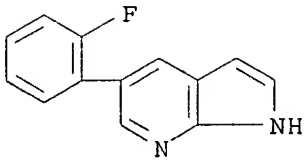
RN 611205-10-8 CAPLUS
CN 1H-Pyrrolo[2,3-b]pyridine, 5-(3-thienyl)- (9CI) (CA INDEX NAME)



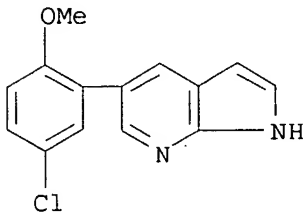
RN 611205-11-9 CAPLUS
CN 1H-Pyrrolo[2,3-b]pyridine, 5-(4-methoxyphenyl)- (9CI) (CA INDEX NAME)



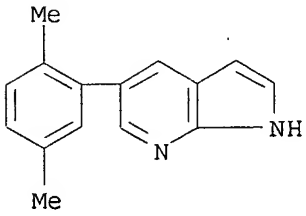
RN 611205-12-0 CAPLUS
CN 1H-Pyrrolo[2,3-b]pyridine, 5-(2-fluorophenyl)- (9CI) (CA INDEX NAME)



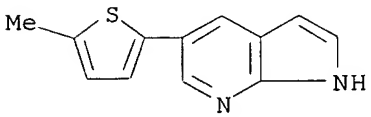
RN 611205-13-1 CAPLUS
CN 1H-Pyrrolo[2,3-b]pyridine, 5-(5-chloro-2-methoxyphenyl)- (9CI) (CA INDEX NAME)



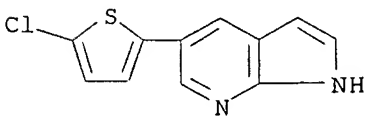
RN 611205-14-2 CAPLUS
CN 1H-Pyrrolo[2,3-b]pyridine, 5-(2,5-dimethylphenyl)- (9CI) (CA INDEX NAME)



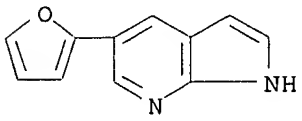
RN 611205-15-3 CAPLUS
CN 1H-Pyrrolo[2,3-b]pyridine, 5-(5-methyl-2-thienyl)- (9CI) (CA INDEX NAME)



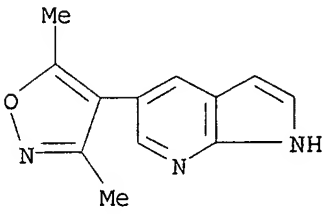
RN 611205-16-4 CAPLUS
CN 1H-Pyrrolo[2,3-b]pyridine, 5-(5-chloro-2-thienyl)- (9CI) (CA INDEX NAME)



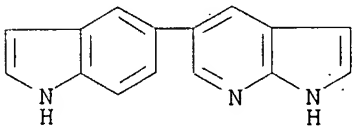
RN 611205-17-5 CAPLUS
CN 1H-Pyrrolo[2,3-b]pyridine, 5-(2-furanyl)- (9CI) (CA INDEX NAME)



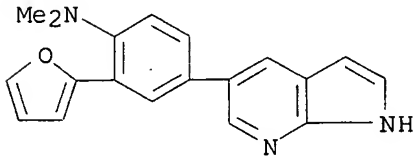
RN 611205-18-6 CAPLUS
CN 1H-Pyrrolo[2,3-b]pyridine, 5-(3,5-dimethyl-4-isoxazolyl)- (9CI) (CA INDEX NAME)



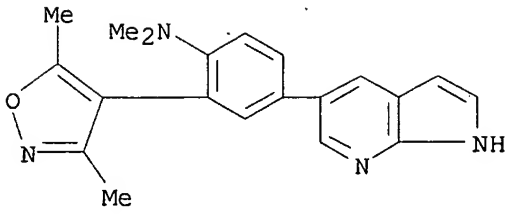
RN 611205-19-7 CAPLUS
CN 1H-Pyrrolo[2,3-b]pyridine, 5-(1H-indol-5-yl)- (9CI) (CA INDEX NAME)



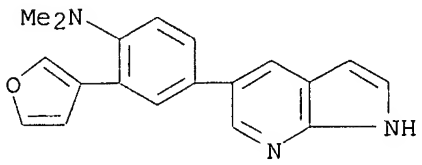
RN 611205-20-0 CAPLUS
CN Benzenamine, 2-(2-furanyl)-N,N-dimethyl-4-(1H-pyrrolo[2,3-b]pyridin-5-yl)- (9CI) (CA INDEX NAME)



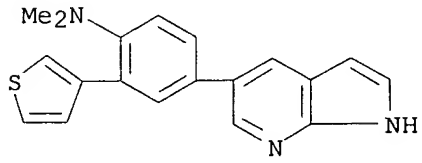
RN 611205-21-1 CAPLUS
CN Benzenamine, 2-(3,5-dimethyl-4-isoxazolyl)-N,N-dimethyl-4-(1H-pyrrolo[2,3-b]pyridin-5-yl)- (9CI) (CA INDEX NAME)



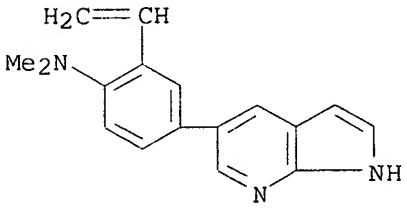
RN 611205-22-2 CAPLUS
CN Benzenamine, 2-(3-furanyl)-N,N-dimethyl-4-(1H-pyrrolo[2,3-b]pyridin-5-yl)-
(9CI) (CA INDEX NAME)



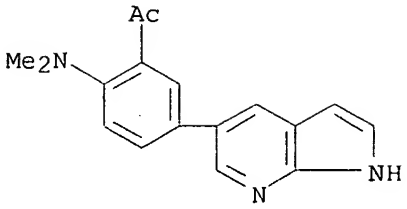
RN 611205-23-3 CAPLUS
CN Benzenamine, N,N-dimethyl-4-(1H-pyrrolo[2,3-b]pyridin-5-yl)-2-(3-thienyl)-
(9CI) (CA INDEX NAME)



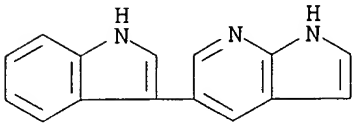
RN 611205-24-4 CAPLUS
CN Benzenamine, 2-ethenyl-N,N-dimethyl-4-(1H-pyrrolo[2,3-b]pyridin-5-yl)-
(9CI) (CA INDEX NAME)



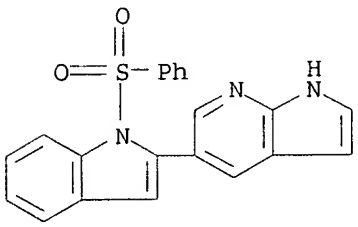
RN 611205-25-5 CAPLUS
CN Ethanone, 1-[2-(dimethylamino)-5-(1H-pyrrolo[2,3-b]pyridin-5-yl)phenyl]-
(9CI) (CA INDEX NAME)



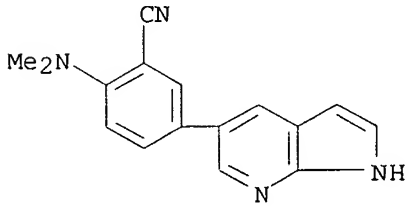
RN 611205-26-6 CAPLUS
CN 1H-Pyrrolo[2,3-b]pyridine, 5-(1H-indol-3-yl)- (9CI) (CA INDEX NAME)



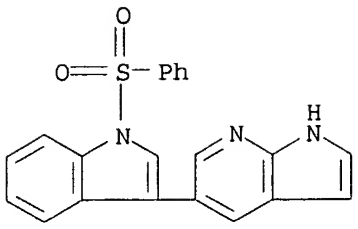
RN 611205-27-7 CAPLUS
CN 1H-Indole, 1-(phenylsulfonyl)-2-(1H-pyrrolo[2,3-b]pyridin-5-yl)- (9CI) (CA INDEX NAME)



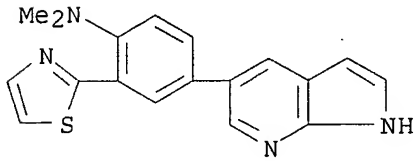
RN 611205-28-8 CAPLUS
CN Benzonitrile, 2-(dimethylamino)-5-(1H-pyrrolo[2,3-b]pyridin-5-yl)- (9CI) (CA INDEX NAME)



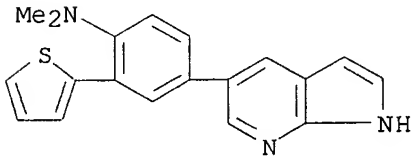
RN 611205-29-9 CAPLUS
CN 1H-Indole, 1-(phenylsulfonyl)-3-(1H-pyrrolo[2,3-b]pyridin-5-yl)- (9CI) (CA INDEX NAME)



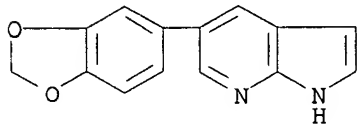
RN 611205-30-2 CAPLUS
CN Benzenamine, N,N-dimethyl-4-(1H-pyrrolo[2,3-b]pyridin-5-yl)-2-(2-thiazolyl)- (9CI) (CA INDEX NAME)



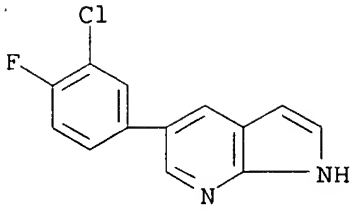
RN 611205-31-3 CAPLUS
CN Benzenamine, N,N-dimethyl-4-(1H-pyrrolo[2,3-b]pyridin-5-yl)-2-(2-thienyl)- (9CI) (CA INDEX NAME)



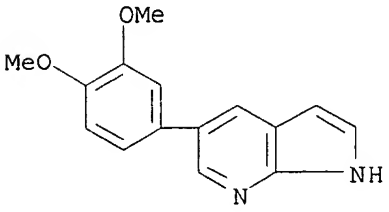
RN 611205-32-4 CAPLUS
CN 1H-Pyrrolo[2,3-b]pyridine, 5-(1,3-benzodioxol-5-yl)- (9CI) (CA INDEX NAME)



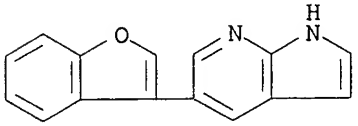
RN 611205-33-5 CAPLUS
CN 1H-Pyrrolo[2,3-b]pyridine, 5-(3-chloro-4-fluorophenyl)- (9CI) (CA INDEX NAME)



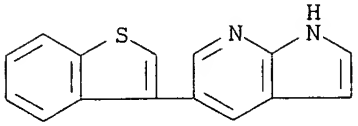
RN 611205-34-6 CAPLUS
CN 1H-Pyrrolo[2,3-b]pyridine, 5-(3,4-dimethoxyphenyl)- (9CI) (CA INDEX NAME)



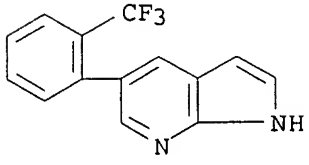
RN 611205-35-7 CAPLUS
CN 1H-Pyrrolo[2,3-b]pyridine, 5-(3-benzofuranyl)- (9CI) (CA INDEX NAME)



RN 611205-36-8 CAPLUS
CN 1H-Pyrrolo[2,3-b]pyridine, 5-benzo[b]thien-3-yl- (9CI) (CA INDEX NAME)

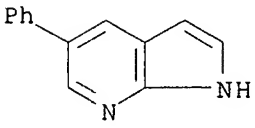


RN 611205-37-9 CAPLUS
CN 1H-Pyrrolo[2,3-b]pyridine, 5-[2-(trifluoromethyl)phenyl]- (9CI) (CA INDEX NAME)

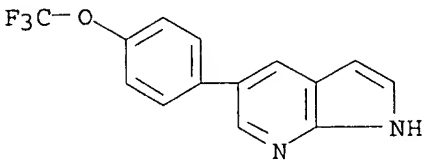


RN 611205-38-0 CAPLUS

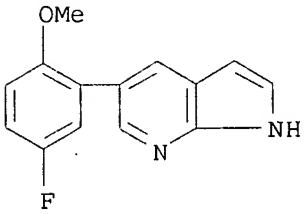
CN 1H-Pyrrolo[2,3-b]pyridine, 5-phenyl- (9CI) (CA INDEX NAME)



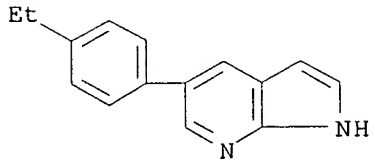
RN 611205-39-1 CAPLUS
CN 1H-Pyrrolo[2,3-b]pyridine, 5-[4-(trifluoromethoxy)phenyl]- (9CI) (CA INDEX NAME)



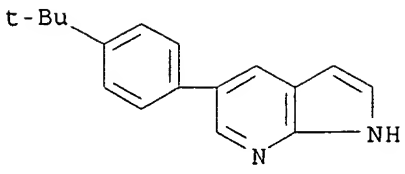
RN 611205-40-4 CAPLUS
CN 1H-Pyrrolo[2,3-b]pyridine, 5-(5-fluoro-2-methoxyphenyl)- (9CI) (CA INDEX NAME)



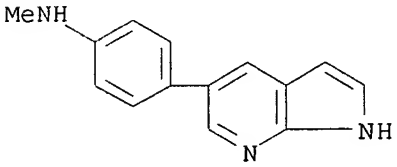
RN 611205-41-5 CAPLUS
CN 1H-Pyrrolo[2,3-b]pyridine, 5-(4-ethylphenyl)- (9CI) (CA INDEX NAME)



RN 611205-42-6 CAPLUS
CN 1H-Pyrrolo[2,3-b]pyridine, 5-[4-(1,1-dimethylethyl)phenyl]- (9CI) (CA INDEX NAME)



RN 611205-43-7 CAPLUS
CN Benzenamine, N-methyl-4-(1H-pyrrolo[2,3-b]pyridin-5-yl)- (9CI) (CA INDEX NAME)



REFERENCE COUNT: 3 THERE ARE 3 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L9 ANSWER 3 OF 8 CAPLUS COPYRIGHT 2006 ACS on STN
ACCESSION NUMBER: 2003:796704 CAPLUS
DOCUMENT NUMBER: 139:307749
TITLE: Preparation of 7-azaindoles as inhibitors of c-Jun N-terminal kinases for treatment of neurodegenerative disorders
INVENTOR(S): Graczyk, Piotr; Numata, Hirotoshi; Khan, Afzal; Palmer, Vanessa
PATENT ASSIGNEE(S): Eisai London Research Laboratories Limited, UK
SOURCE: PCT Int. Appl., 70 pp.
CODEN: PIXXD2
DOCUMENT TYPE: Patent
LANGUAGE: English
FAMILY ACC. NUM. COUNT: 1
PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
WO 2003082868	A1	20031009	WO 2003-GB1112	20030317 <--
W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NI, NO, NZ, OM, PH, PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, YU, ZA, ZM, ZW				
RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM, AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IT, LU, MC, NL, PT, RO, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG				
CA 2480317	AA	20031009	CA 2003-2480317	20030317 <--
AU 2003214412	A1	20031013	AU 2003-214412	20030317 <--
EP 1490364	A1	20041229	EP 2003-709984	20030317
R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, SI, LT, LV, FI, RO, MK, CY, AL, TR, BG, CZ, EE, HU, SK				
CN 1656094	A	20050817	CN 2003-812103	20030317
JP 2005534618	T2	20051117	JP 2003-580333	20030317
US 2005272761	A1	20051208	US 2005-509128	20050728

PRIORITY APPLN. INFO.:

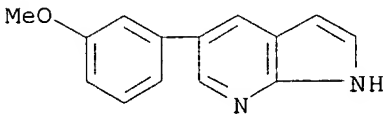
GB 2002-7491	A	20020328
GB 2002-17330	A	20020725
WO 2003-GB1112	W	20030317

OTHER SOURCE(S): MARPAT 139:307749

IT 344454-28-0P 611204-93-4P 611204-95-6P
611204-96-7P 611204-97-8P 611204-98-9P
611204-99-0P 611205-00-6P 611205-01-7P
611205-02-8P 611205-03-9P 611205-04-0P
611205-05-1P 611205-06-2P 611205-07-3P
611205-08-4P 611205-09-5P 611205-10-8P
611205-11-9P 611205-12-0P 611205-13-1P
611205-14-2P
RL: PAC (Pharmacological activity); SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)
(drug candidate; preparation of azaindoles as inhibitors of c-jun N-terminal kinases for treatment of neurodegenerative disorders)

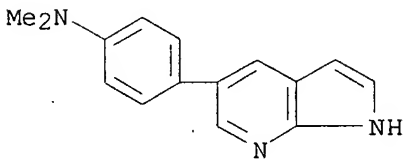
RN 344454-28-0 CAPLUS

CN 1H-Pyrrolo[2,3-b]pyridine, 5-(3-methoxyphenyl)- (9CI) (CA INDEX NAME)



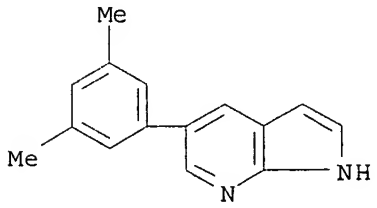
RN 611204-93-4 CAPLUS

CN Benzenamine, N,N-dimethyl-4-(1H-pyrrolo[2,3-b]pyridin-5-yl)- (9CI) (CA INDEX NAME)



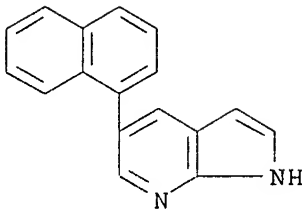
RN 611204-95-6 CAPLUS

CN 1H-Pyrrolo[2,3-b]pyridine, 5-(3,5-dimethylphenyl)- (9CI) (CA INDEX NAME)

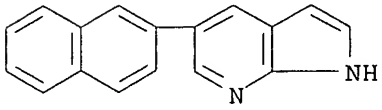


RN 611204-96-7 CAPLUS

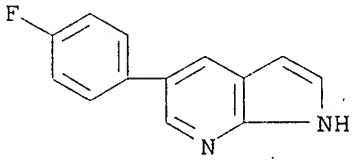
CN 1H-Pyrrolo[2,3-b]pyridine, 5-(1-naphthalenyl)- (9CI) (CA INDEX NAME)



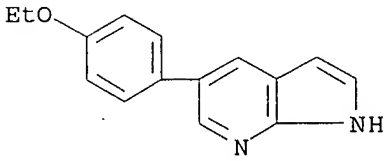
RN 611204-97-8 CAPLUS
CN 1H-Pyrrolo[2,3-b]pyridine, 5-(2-naphthalenyl)- (9CI) (CA INDEX NAME)



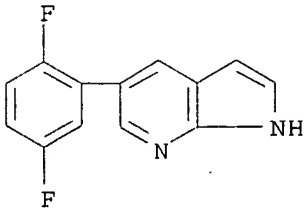
RN 611204-98-9 CAPLUS
CN 1H-Pyrrolo[2,3-b]pyridine, 5-(4-fluorophenyl)- (9CI) (CA INDEX NAME)



RN 611204-99-0 CAPLUS
CN 1H-Pyrrolo[2,3-b]pyridine, 5-(4-ethoxyphenyl)- (9CI) (CA INDEX NAME)

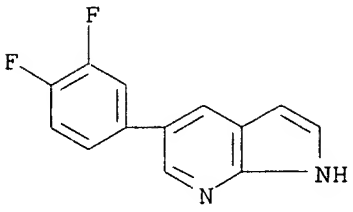


RN 611205-00-6 CAPLUS
CN 1H-Pyrrolo[2,3-b]pyridine, 5-(2,5-difluorophenyl)- (9CI) (CA INDEX NAME)

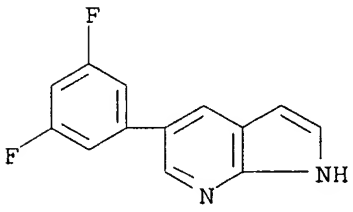


RN 611205-01-7 CAPLUS

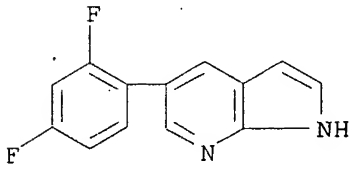
CN 1H-Pyrrolo[2,3-b]pyridine, 5-(3,4-difluorophenyl)- (9CI) (CA INDEX NAME)



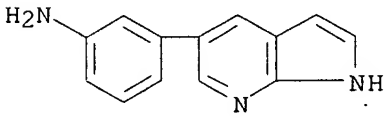
RN 611205-02-8 CAPLUS
CN 1H-Pyrrolo[2,3-b]pyridine, 5-(3,5-difluorophenyl)- (9CI) (CA INDEX NAME)



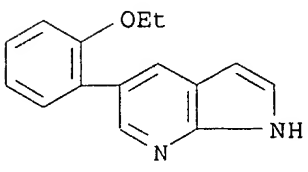
RN 611205-03-9 CAPLUS
CN 1H-Pyrrolo[2,3-b]pyridine, 5-(2,4-difluorophenyl)- (9CI) (CA INDEX NAME)



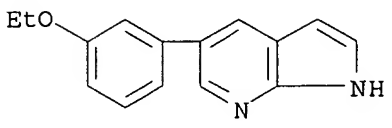
RN 611205-04-0 CAPLUS
CN Benzenamine, 3-(1H-pyrrolo[2,3-b]pyridin-5-yl)- (9CI) (CA INDEX NAME)



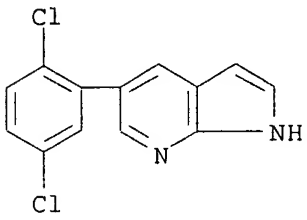
RN 611205-05-1 CAPLUS
CN 1H-Pyrrolo[2,3-b]pyridine, 5-(2-ethoxyphenyl)- (9CI) (CA INDEX NAME)



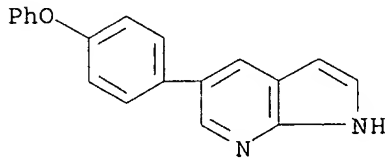
RN 611205-06-2 CAPLUS
CN 1H-Pyrrolo[2,3-b]pyridine, 5-(3-ethoxyphenyl)- (9CI) (CA INDEX NAME)



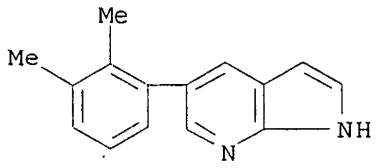
RN 611205-07-3 CAPLUS
CN 1H-Pyrrolo[2,3-b]pyridine, 5-(2,5-dichlorophenyl)- (9CI) (CA INDEX NAME)



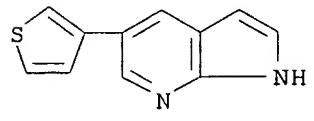
RN 611205-08-4 CAPLUS
CN 1H-Pyrrolo[2,3-b]pyridine, 5-(4-phenoxyphenyl)- (9CI) (CA INDEX NAME)



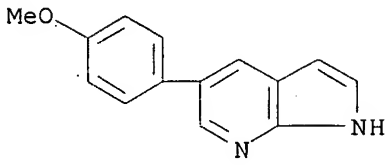
RN 611205-09-5 CAPLUS
CN 1H-Pyrrolo[2,3-b]pyridine, 5-(2,3-dimethylphenyl)- (9CI) (CA INDEX NAME)



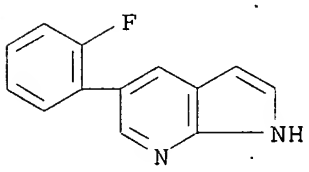
RN 611205-10-8 CAPLUS
CN 1H-Pyrrolo[2,3-b]pyridine, 5-(3-thienyl)- (9CI) (CA INDEX NAME)



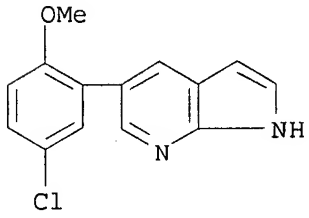
RN 611205-11-9 CAPLUS
CN 1H-Pyrrolo[2,3-b]pyridine, 5-(4-methoxyphenyl)- (9CI) (CA INDEX NAME)



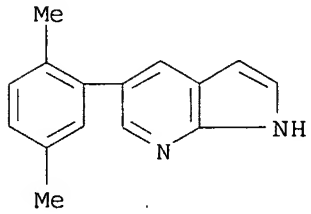
RN 611205-12-0 CAPLUS
CN 1H-Pyrrolo[2,3-b]pyridine, 5-(2-fluorophenyl)- (9CI) (CA INDEX NAME)



RN 611205-13-1 CAPLUS
CN 1H-Pyrrolo[2,3-b]pyridine, 5-(5-chloro-2-methoxyphenyl)- (9CI) (CA INDEX NAME)

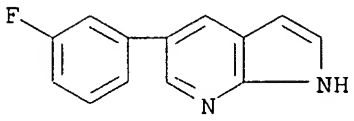


RN 611205-14-2 CAPLUS
CN 1H-Pyrrolo[2,3-b]pyridine, 5-(2,5-dimethylphenyl)- (9CI) (CA INDEX NAME)



IT 611204-92-3P
RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation); RACT (Reactant or reagent)
(intermediate; preparation of azaindoles as inhibitors of c-jun N-terminal kinases for treatment of neurodegenerative disorders)

RN 611204-92-3 CAPLUS
CN 1H-Pyrrolo[2,3-b]pyridine, 5-(3-fluorophenyl)- (9CI) (CA INDEX NAME)



REFERENCE COUNT: 4 THERE ARE 4 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L9 ANSWER 4 OF 8 CAPLUS COPYRIGHT 2006 ACS on STN
ACCESSION NUMBER: 2003:777399 CAPLUS
DOCUMENT NUMBER: 139:292151
TITLE: Preparation of pyridine derivatives as protein kinase inhibitors
INVENTOR(S): Li, Qun; Woods, Keith W.; Zhu, Gui-Dong; Fischer, John P.; Gong, Jianchun; Li, Tongmei; Gandhi, Virajkumar; Thomas, Sheela A.; Packard, Garrick K.; Song, Xiaohong; Abrams, Jason N.; Diebold, Robert; Dinges, Jurgen; Hutchins, Charles; Stoll, Vincent S.; Rosenberg, Saul H.; Giranda, Vincent L.
PATENT ASSIGNEE(S): USA
SOURCE: U.S. Pat. Appl. Publ., 120 pp., Cont.-in-part of U.S. Ser. No. 23,363, abandoned.
CODEN: USXXCO
DOCUMENT TYPE: Patent
LANGUAGE: English
FAMILY ACC. NUM. COUNT: 2
PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
US 2003187026	A1	20031002	US 2002-295833	20021118 <--
CA 2470214	AA	20030626	CA 2002-2470214	20021212 <--
WO 2003051366	A2	20030626	WO 2002-US39915	20021212 <--
WO 2003051366	A3	20040325		
W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, OM, PH, PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, TJ, TM, TN, TR, TT, TZ, UA, UG, UZ, VC, VN, YU, ZA, ZM, ZW				
RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM, AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG				
AU 2002353147	A1	20030630	AU 2002-353147	20021212 <--
EP 1463505	A2	20041006	EP 2002-790126	20021212
R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, SI, LT, LV, FI, RO, MK, CY, AL, TR, BG, CZ, EE, SK				
JP 2005516927	T2	20050609	JP 2003-552299	20021212
PRIORITY APPLN. INFO.:			US 2001-23363	B2 20011213
			US 2002-295833	A 20021118
			WO 2002-US39915	W 20021212

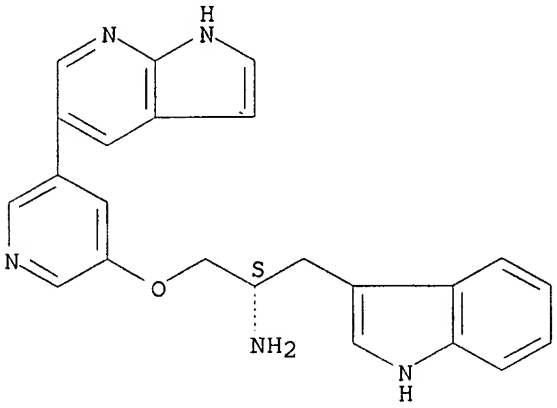
OTHER SOURCE(S): MARPAT 139:292151
IT 552326-49-5P 552326-50-8P
RL: PAC (Pharmacological activity); SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)

(preparation of pyridine derivs. as protein kinase inhibitors)

RN 552326-49-5 CAPLUS

CN 1H-Indole-3-ethanamine, α-[[[5-(1H-pyrrolo[2,3-b]pyridin-5-yl)-3-pyridinyl]oxy]methyl]-, (αS)- (9CI) (CA INDEX NAME)

Absolute stereochemistry.



RN 552326-50-8 CAPLUS

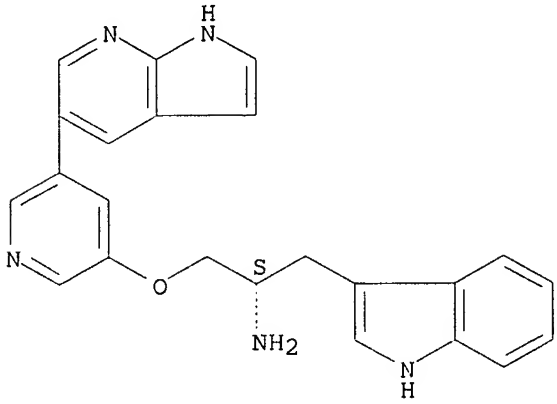
CN 1H-Indole-3-ethanamine, α-[[[5-(1H-pyrrolo[2,3-b]pyridin-5-yl)-3-pyridinyl]oxy]methyl]-, (αS)-, trifluoroacetate (9CI) (CA INDEX NAME)

CM 1

CRN 552326-49-5

CMF C23 H21 N5 O

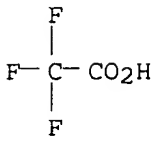
Absolute stereochemistry.



CM 2

CRN 76-05-1

CMF C2 H F3 O2



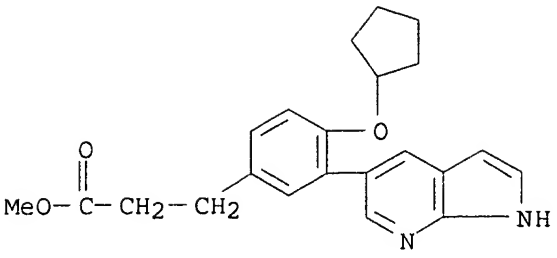
L9 ANSWER 5 OF 8 CAPLUS COPYRIGHT 2006 ACS on STN
ACCESSION NUMBER: 2003:678772 CAPLUS
DOCUMENT NUMBER: 139:214465
TITLE: Preparation of substituted phenylalkanoic acid derivatives as inhibitors of prostaglandin and leukotriene production
INVENTOR(S): Shoda, Motoshi; Kuriyama, Hiroshi
PATENT ASSIGNEE(S): Asahi Kasei Kabushiki Kaisha, Japan
SOURCE: PCT Int. Appl., 607 pp.
CODEN: PIXXD2
DOCUMENT TYPE: Patent
LANGUAGE: Japanese
FAMILY ACC. NUM. COUNT: 1
PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
WO 2003070686	A1	20030828	WO 2003-JP1849	20030220 <--
W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, OM, PH, PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, YU, ZA, ZM, ZW				
RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM, AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IT, LU, MC, NL, PT, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG				
CA 2477208	AA	20030828	CA 2003-2477208	20030220 <--
AU 2003211384	A1	20030909	AU 2003-211384	20030220 <--
US 2004044258	A1	20040304	US 2003-368435	20030220
US 6867320	B2	20050315		
EP 1477472	A1	20041117	EP 2003-706983	20030220
R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, SI, LT, LV, FI, RO, MK, CY, AL, TR, BG, CZ, EE, HU, SK				
CN 1653032	A	20050810	CN 2003-808999	20030220

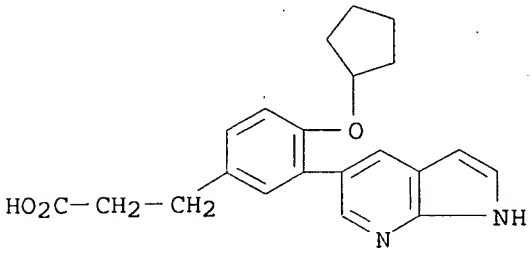
PRIORITY APPLN. INFO.:

JP 2002-45293	A	20020221
JP 2002-301543	A	20021016
US 2002-358337P	P	20020222
US 2002-419098P	P	20021018
WO 2003-JP1849	W	20030220

OTHER SOURCE(S): MARPAT 139:214465
IT 590416-03-8P 590416-04-9P
RL: PAC (Pharmacological activity); SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)
(preparation of substituted phenylalkanoic acid derivs. as inhibitors of prostaglandin and leukotriene production for prevention or treatment of inflammations, allergies, and autoimmune diseases, and for antipyresis and/or analgesia)
RN 590416-03-8 CAPLUS
CN Benzenepropanoic acid, 4-(cyclopentyloxy)-3-(1H-pyrrolo[2,3-b]pyridin-5-yl)-, methyl ester (9CI) (CA INDEX NAME)



RN 590416-04-9 CAPLUS
CN Benzenepropanoic acid, 4-(cyclopentyloxy)-3-(1H-pyrrolo[2,3-b]pyridin-5-yl)- (9CI) (CA INDEX NAME)



REFERENCE COUNT: 5 THERE ARE 5 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L9 ANSWER 6 OF 8 CAPLUS COPYRIGHT 2006 ACS on STN
ACCESSION NUMBER: 2003:491046 CAPLUS
DOCUMENT NUMBER: 139:69152
TITLE: Preparation of pyridine derivatives as protein kinase inhibitors
INVENTOR(S): Li, Qun; Woods, Keith W.; Zhu, Gui-Dong; Fischer, John P.; Gong, Jianchun; Li, Tongmei; Gandhi, Viraj; Thomas, Sheela A.; Packard, Garrick; Song, Xiaohong; Abrams, Jason N.; Diebold, Robert; Dinges, Jurgen; Hutchins, Charles; Stoll, Vincent S.; Rosenberg, Saul H.; Giranda, Vincent L.
PATENT ASSIGNEE(S): Abbott Laboratories, USA
SOURCE: PCT Int. Appl., 261 pp.
CODEN: PIXXD2
DOCUMENT TYPE: Patent
LANGUAGE: English
FAMILY ACC. NUM. COUNT: 2
PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
WO 2003051366	A2	20030626	WO 2002-US39915	20021212 <--
WO 2003051366	A3	20040325		

W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, OM, PH, PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, TJ, TM, TN, TR, TT, TZ, UA, UG, UZ, VC, VN, YU, ZA, ZM, ZW
RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ, BY,

KG, KZ, MD, RU, TJ, TM, AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG

US 2003187026	A1	20031002	US 2002-295833		20021118 <--
CA 2470214	AA	20030626	CA 2002-2470214		20021212 <--
AU 2002353147	A1	20030630	AU 2002-353147		20021212 <--
EP 1463505	A2	20041006	EP 2002-790126		20021212
R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, SI, LT, LV, FI, RO, MK, CY, AL, TR, BG, CZ, EE, SK					
JP 2005516927	T2	20050609	JP 2003-552299		20021212
PRIORITY APPLN. INFO.:			US 2001-23363	A	20011213
			US 2002-295833	A	20021118
			WO 2002-US39915	W	20021212

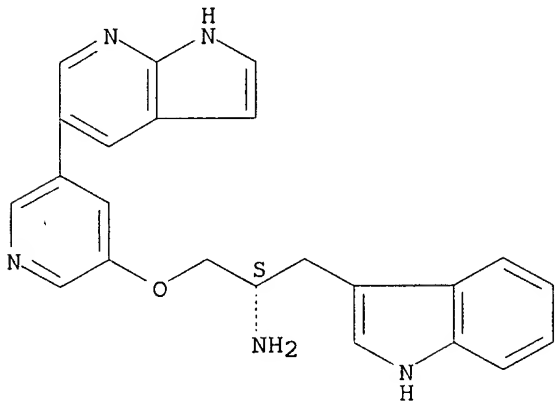
OTHER SOURCE(S): MARPAT 139:69152

IT 552326-49-5P 552326-50-8P
RL: SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)
(preparation of pyridine derivs. as protein kinase inhibitors)

RN 552326-49-5 CAPLUS

CN 1H-Indole-3-ethanamine, α-[[[5-(1H-pyrrolo[2,3-b]pyridin-5-yl)-3-pyridinyl]oxy]methyl]-, (αS)- (9CI) (CA INDEX NAME)

Absolute stereochemistry.



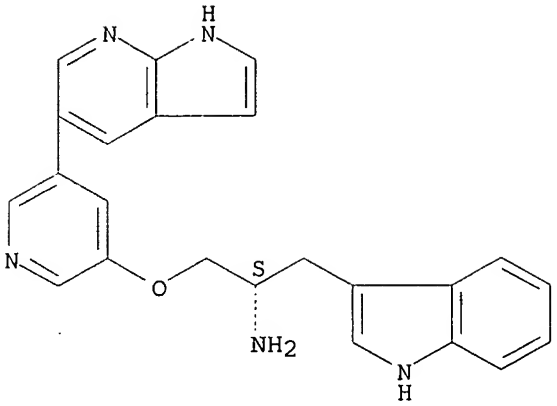
RN 552326-50-8 CAPLUS

CN 1H-Indole-3-ethanamine, α-[[[5-(1H-pyrrolo[2,3-b]pyridin-5-yl)-3-pyridinyl]oxy]methyl]-, (αS)-, trifluoroacetate (9CI) (CA INDEX NAME)

CM 1

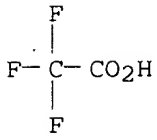
CRN 552326-49-5
CMF C23 H21 N5 O

Absolute stereochemistry.



CM 2

CRN 76-05-1
CMF C2 H F3 O2

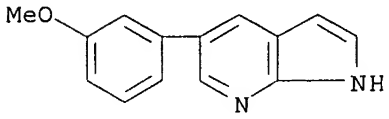


L9 ANSWER 7 OF 8 CAPLUS COPYRIGHT 2006 ACS on STN
ACCESSION NUMBER: 2001:507532 CAPLUS
DOCUMENT NUMBER: 135:107148
TITLE: Preparation of N-cyanomethyl amides as cysteine protease inhibitors
INVENTOR(S): Oballa, Renata Marcella; Prasit, Petpiboon; Robichaud, Joel Stephane; Isabel, Elise; Mendonca, Rohan V.; Venkatraman, Shankar; Setti, Eduardo; Wang, Dan-Xiong
PATENT ASSIGNEE(S): Merck Frosst Canada & Co., Can.; Axys Pharmaceuticals, Inc.
SOURCE: PCT Int. Appl., 157 pp.
CODEN: PIXXD2
DOCUMENT TYPE: Patent
LANGUAGE: English
FAMILY ACC. NUM. COUNT: 1
PATENT INFORMATION:

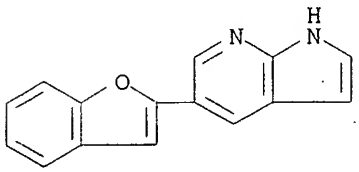
PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
WO 2001049288	A1	20010712	WO 2001-US341	20010105 <--
W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CR, CU, CZ, DE, DK, DM, DZ, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, TZ, UA, UG, US, UZ, VN, YU, ZA, ZW				
RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZW, AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG				
CA 2396257	AA	20010712	CA 2001-2396257	20010105 <--

IT 344454-28-0 344454-31-5 344454-45-1
RL: RCT (Reactant); RACT (Reactant or reagent)
(synthesis of fluorescent substances and application for obtaining fluorescence probes and detection of PCR products)

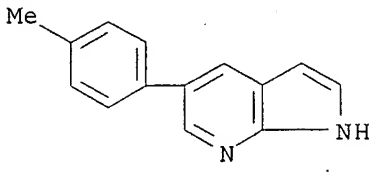
RN 344454-28-0 CAPLUS
CN 1H-Pyrrolo[2,3-b]pyridine, 5-(3-methoxyphenyl)- (9CI) (CA INDEX NAME)



RN 344454-31-5 CAPLUS
CN 1H-Pyrrolo[2,3-b]pyridine, 5-(2-benzofuranyl)- (9CI) (CA INDEX NAME)



RN 344454-45-1 CAPLUS
CN 1H-Pyrrolo[2,3-b]pyridine, 5-(4-methylphenyl)- (9CI) (CA INDEX NAME)



US 2002052378	A1	20020502	US 2001-754962		20010105 <--
US 6525036	B2	20030225			
EP 1248612	A1	20021016	EP 2001-900903		20010105 <--
R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, SI, LT, LV, FI, RO, MK, CY, AL, TR					
JP 2003525874	T2	20030902	JP 2001-549656		20010105 <--
AU 779855	B2	20050217	AU 2001-26314		20010105
PRIORITY APPLN. INFO.:			US 2000-174978P	P	20000106
			US 2000-256793P	P	20001219
			WO 2001-US341	W	20010105

OTHER SOURCE(S): MARPAT 135:107148

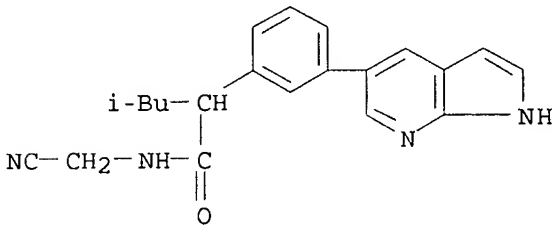
IT 349669-75-6P

RL: BAC (Biological activity or effector, except adverse); BSU (Biological study, unclassified); SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)

(preparation of N-cyanomethyl amides as protease cysteine inhibitors)

RN 349669-75-6 CAPLUS

CN Benzeneacetamide, N-(cyanomethyl)-α-(2-methylpropyl)-3-(1H-pyrrolo[2,3-b]pyridin-5-yl)- (9CI) (CA INDEX NAME)



L9 ANSWER 8 OF 8 CAPLUS COPYRIGHT 2006 ACS on STN

ACCESSION NUMBER: 2001:432896 CAPLUS

DOCUMENT NUMBER: 135:43132

TITLE: Synthesis of fluorescent substances and application for obtaining fluorescence probes and detection of PCR products

INVENTOR(S): Inomata, Hiroko; Shinoki, Hiroshi; Kojima, Masayoshi; Sudo, Yukio; Nishigaki, Junji; Seshimoto, Osamu

PATENT ASSIGNEE(S): Fuji Photo Film Co., Ltd., Japan

SOURCE: Eur. Pat. Appl., 41 pp.
CODEN: EPXXDW

DOCUMENT TYPE: Patent

LANGUAGE: English

FAMILY ACC. NUM. COUNT: 1

PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.		DATE
EP 1106621	A2	20010613	EP 2000-126447		20001206 <--
EP 1106621	A3	20010912			
EP 1106621	B1	20031119			
R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, SI, LT, LV, FI, RO					
JP 2001163895	A2	20010619	JP 1999-347886		19991207 <--
JP 2001163900	A2	20010619	JP 1999-348015		19991207 <--
US 2003013088	A1	20030116	US 2000-731279		20001206 <--
US 6642375	B2	20031104			
PRIORITY APPLN. INFO.:			JP 1999-347886	A	19991207
			JP 1999-348015	A	19991207

OTHER SOURCE(S): MARPAT 135:43132